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FILE COVERS 1907 - 20 Oct 2008 VOL 149 ISS 17
FILE LAST UPDATED: 19 Oct 2008 (20081019/ED)

Caplus now includes complete International Patent Classification (IPC) reclassification data for the second quarter of 2008.

Effective October 17, 2005, revised CAS Information Use Policies apply. They are available for your review at:

<http://www.cas.org/legal/infopolicy.html>

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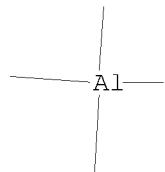
Uploading C:\Program Files\Stnexp\Queries\10642438.str

L1 STRUCTURE UPLOADED

=> d

L1 HAS NO ANSWERS

L1 STR



Structure attributes must be viewed using STN Express query preparation.

=> s l1 and quaternary?

REGISTRY INITIATED

Substance data SEARCH and crossover from CAS REGISTRY in progress...

Use DISPLAY HITSTR (or FHITSTR) to directly view retrieved structures.

SAMPLE SEARCH INITIATED 14:06:29 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED - 1747 TO ITERATE

100.0% PROCESSED 1747 ITERATIONS
SEARCH TIME: 00.00.01

34 ANSWERS

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
BATCH **COMPLETE**

TOh

20/10/2008

10/923,271

PROJECTED ITERATIONS: 32433 TO 37447
PROJECTED ANSWERS: 331 TO 1029

L2 34 SEA SSS SAM L1

L3 42 L2

142881 QUATERNARY?
L4 0 L3 AND QUATERNARY?

=> s l1 and docusate
REGISTRY INITIATED
Substance data SEARCH and crossover from CAS REGISTRY in progress...
Use DISPLAY HITSTR (or FHITSTR) to directly view retrieved structures.

SAMPLE SEARCH INITIATED 14:07:12 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED - 1747 TO ITERATE

100.0% PROCESSED 1747 ITERATIONS 34 ANSWERS
SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
BATCH **COMPLETE**
PROJECTED ITERATIONS: 32433 TO 37447
PROJECTED ANSWERS: 331 TO 1029

L5 34 SEA SSS SAM L1

L6 42 L5

335 DOCUSATE
L7 0 L6 AND DOCUSATE

=> s l1 and ionic liquid
REGISTRY INITIATED
Substance data SEARCH and crossover from CAS REGISTRY in progress...
Use DISPLAY HITSTR (or FHITSTR) to directly view retrieved structures.

SAMPLE SEARCH INITIATED 14:07:41 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED - 1747 TO ITERATE

100.0% PROCESSED 1747 ITERATIONS 34 ANSWERS

10/923,271

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
BATCH **COMPLETE**
PROJECTED ITERATIONS: 32433 TO 37447
PROJECTED ANSWERS: 331 TO 1029

L8 34 SEA SSS SAM L1

L9 42 L8

298217 IONIC
4 LIQUID
0 IONIC LIQUID
(IONIC(W)LIQUID)
L10 0 L9 AND IONIC LIQUID

=> s l1 and ionic liquid

REGISTRY INITIATED

Substance data SEARCH and crossover from CAS REGISTRY in progress...
Use DISPLAY HITSTR (or FHITSTR) to directly view retrieved structures.

SAMPLE SEARCH INITIATED 14:07:57 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED - 1747 TO ITERATE

100.0% PROCESSED 1747 ITERATIONS 34 ANSWERS
SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
BATCH **COMPLETE**
PROJECTED ITERATIONS: 32433 TO 37447
PROJECTED ANSWERS: 331 TO 1029

L11 34 SEA SSS SAM L1

L12 42 L11

298217 IONIC
850264 LIQUID
4497 IONIC LIQUID
(IONIC(W)LIQUID)
L13 0 L12 AND IONIC LIQUID

=> s l1 and ionic compo?

REGISTRY INITIATED

Substance data SEARCH and crossover from CAS REGISTRY in progress...

10/923,271

Use DISPLAY HITSTR (or FHITSTR) to directly view retrieved structures.

SAMPLE SEARCH INITIATED 14:08:20 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED - 1747 TO ITERATE

100.0% PROCESSED 1747 ITERATIONS 34 ANSWERS
SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
BATCH **COMPLETE**
PROJECTED ITERATIONS: 32433 TO 37447
PROJECTED ANSWERS: 331 TO 1029

L14 34 SEA SSS SAM L1

L15 42 L14

298217 IONIC
4073012 COMPO?
3077 IONIC COMPO?
(IONIC(W)COMPO?)

L16 0 L15 AND IONIC COMPO?

=> s l1 and compo?

REGISTRY INITIATED

Substance data SEARCH and crossover from CAS REGISTRY in progress...
Use DISPLAY HITSTR (or FHITSTR) to directly view retrieved structures.

SAMPLE SEARCH INITIATED 14:08:57 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED - 1747 TO ITERATE

100.0% PROCESSED 1747 ITERATIONS 34 ANSWERS
SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
BATCH **COMPLETE**
PROJECTED ITERATIONS: 32433 TO 37447
PROJECTED ANSWERS: 331 TO 1029

L17 34 SEA SSS SAM L1

L18 42 L17

10/923,271

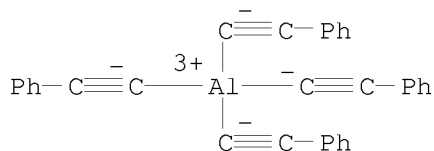
4073012 COMPO?
L19 12 L18 AND COMPO?

=> s l19 and ammonium
431907 AMMONIUM
L20 0 L19 AND AMMONIUM

=> s l19 and py<2002
21968514 PY<2002
L21 10 L19 AND PY<2002

=> d 1-10 ibib abs hitstr

L21 ANSWER 1 OF 10 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 1999:367958 CAPLUS
DOCUMENT NUMBER: 131:185603
TITLE: A novel synthesis of a highly heat-resistant
organosilicon polymer using base catalysts
AUTHOR(S): Itoh, Masayoshi
CORPORATE SOURCE: Organic Performance Materials Laboratory, Mitsui
Chemicals, Inc., Yokohama-city, 247-8567, Japan
SOURCE: Catalysis Surveys from Japan (1999), 3(1),
61-69
CODEN: CSURFY; ISSN: 1384-6574
PUBLISHER: Baltzer Science Publishers
DOCUMENT TYPE: Journal
LANGUAGE: English
AB A new highly heat-resistant polymer containing silicon,
poly[(phenylsilylene)ethynylene-1,3-phenyleneethynylene] (MSP), was prepared
by dehydrogenative coupling polymerization between phenylsilane and
1,3-diethynylbenzene in the presence of base catalysts such as alkaline earth
metal oxides, metal hydrides and metal alkoxides. The preparation process,
catalytic activities, reaction mechanisms and polymer properties were
discussed.
IT 4015-69-4
RL: CAT (Catalyst use); USES (Uses)
(preparation of a highly heat-resistant 1,3-diethynylbenzene-phenylsilane
copolymer using base catalysts)
RN 4015-69-4 CAPLUS
CN Aluminate(1-), tetrakis(phenylethynyl)-, lithium, (T-4)- (9CI) (CA INDEX
NAME)



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REFERENCE COUNT: 18 THERE ARE 18 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L21 ANSWER 2 OF 10 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 1992:266214 CAPLUS
DOCUMENT NUMBER: 116:266214
ORIGINAL REFERENCE NO.: 116:44943a,44946a
TITLE: Methods and compounds for forming alkaline earth metal-containing films
INVENTOR(S): Kruck, Thomas; Heck, Stephan
PATENT ASSIGNEE(S): Kali-Chemie A.-G., Germany
SOURCE: Ger. Offen., 11 pp.
CODEN: GWXXBX
DOCUMENT TYPE: Patent
LANGUAGE: German
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
DE 4121369	A1	19920109	DE 1991-4121369	19910628 <--
PRIORITY APPLN. INFO.:			DE 1990-4020976	A1 19900703

OTHER SOURCE(S): MARPAT 116:266214

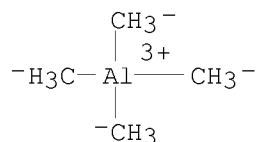
AB The title methods entail the decomposition of compds. described by the general formulas $M(ZR_{14})_2$ (I), $M(ZR_{13}H)_2$ (II), or $M(ZR_{12}H_2)_2$ (III) ($M = Ca, Sr, or Ba, Z = Al, Y, or Sc$, and $R_1 = a$ linear or branched C1-4 alkyl group or an aryl group, especially a Ph group). The compds. may be applied to a substrate as liqs. or vapors. Selected compds. of those described by the formulas I, II, and III are claimed.

IT 141646-37-9P

RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation and use of, in alkaline earth metal-containing film formation)

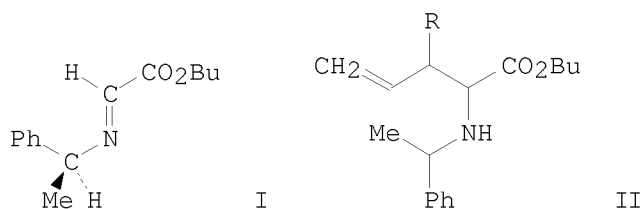
RN 141646-37-9 CAPLUS

CN Aluminate(1-), tetramethyl-, strontium (2:1), (T-4)- (9CI) (CA INDEX NAME)



L21 ANSWER 3 OF 10 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 1989:193346 CAPLUS
DOCUMENT NUMBER: 110:193346
ORIGINAL REFERENCE NO.: 110:32125a,32128a
TITLE: Studies on the reaction of α -imino esters with organometallic compounds

AUTHOR(S): Yamamoto, Yoshinori; Ito, Wataru
 CORPORATE SOURCE: Fac. Sci., Tohoku Univ., Sendai, 980, Japan
 SOURCE: Tetrahedron (1988), 44(17), 5415-23
 CODEN: TETRAB; ISSN: 0040-4020
 DOCUMENT TYPE: Journal
 LANGUAGE: English
 OTHER SOURCE(S): CASREACT 110:193346
 GI



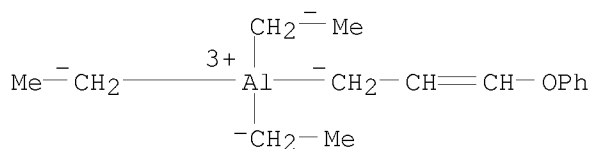
AB Benzylzinc reagent reacted with α -imino ester I at the α -carbon exclusively, though other organometallic reagents, such as Mg, Al, Cu, Ti, and B derivs., reacted at the nitrogen atom. Use of the (S)-amine as a chiral auxiliary of I created the R chirality at the imino carbon. Very high chiral induction was realized in the reaction of prenylzinc reagent with 8-(-)-phenylmethyl N-(methoxyimino)acetate. The reaction of I with heteroatom-substituted allylic organometallic compds. $\text{RCH:CHCH}_2\text{MLn}$ [R = OMe, MLn = ZnBr, Ti(OCHMe₂)₃, AlEt₃Li; R = OPh, MLn = ZnBr, AlEt₃Li; R = SMe, MLn = ZnBr, B(OMe)₂] gave the corresponding α -heteroatom substituted amino acid derivs. II. Here again, the allylic zinc reagent gave the adduct in higher yield than the corresponding Ti, Al, and B reagents.

IT 120169-59-7

RL: RCT (Reactant); RACT (Reactant or reagent)
 (reaction of, with chiral imino ester, stereochem. of)

RN 120169-59-7 CAPLUS

CN Aluminate(1-), triethyl(3-phenoxy-2-propenyl)-, lithium, (T-4)- (9CI) (CA INDEX NAME)



L21 ANSWER 4 OF 10 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 1984:86283 CAPLUS
 DOCUMENT NUMBER: 100:86283

10/923,271

ORIGINAL REFERENCE NO.: 100:13095a,13098a
TITLE: Composition containing chlorine, bromine,
and magnesium suitable as a polymerization catalyst
support
PATENT ASSIGNEE(S): Gulf Research and Development Co. , USA
SOURCE: Neth. Appl., 19 pp.
CODEN: NAXXAN
DOCUMENT TYPE: Patent
LANGUAGE: Dutch
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
NL 8201563	A	19831101	NL 1982-1563	19820414 <--

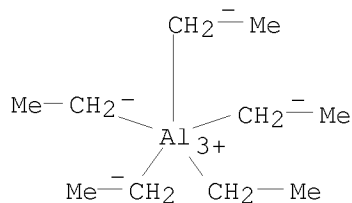
PRIORITY APPLN. INFO.: NL 1982-1563 19820414

AB A catalyst support consists of MgCl₂ doped with Br in a mol. ratio from 1:99 to 50:50 (and especially from 2.5:97.5 to 15:85). Thus, the reaction product of 0.03 mol MgAl₂Et₈ [15415-18-6], 0.056 mmol Et₂AlCl [96-10-6], and 0.002 mol AlBr₃ was further treated with 0.3 mL Et benzoate [93-89-0] and subsequently with TiCl₄ to obtain a catalyst containing Mg 20.7, Al 0.05, Ti 0.8, Br 9.7, and Cl 53.2 weight%. In the polymerization of propene, the catalyst had an activity of 112,500 g polymer/g Ti, and the polymer [25085-53-4] had isotacticity 97% and intrinsic viscosity 3.7 dL/g (ASTM D-2857).

IT 82404-69-1
RL: USES (Uses)
(catalyst compns. containing, for stereospecific polymerization of alkenes)

RN 82404-69-1 CAPLUS

CN Aluminate(2-), pentaethyl-, magnesium (1:1) (CA INDEX NAME)



L21 ANSWER 5 OF 10 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 1983:216200 CAPLUS
DOCUMENT NUMBER: 98:216200
ORIGINAL REFERENCE NO.: 98:32893a,32896a
TITLE: Composition containing chlorine, bromine and
magnesium
INVENTOR(S): Beach, David L.; Zambelli, Adolfo
PATENT ASSIGNEE(S): Gulf Research and Development Co. , USA
SOURCE: U.S., 8 pp.

10/923,271

CODEN: USXXAM
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 4366086	A	19821228	US 1980-221064	19801229 <--
JP 58183707	A	19831027	JP 1982-56777	19820407 <--

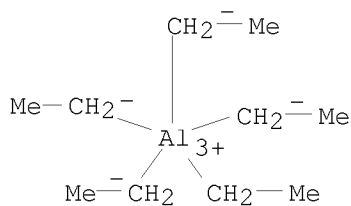
PRIORITY APPLN. INFO.: US 1980-221064 19801229

AB A support for Ziegler catalysts exhibiting high polymer yields and a high degree of stereospecificity is obtained by treating an organo Mg compound with a mixture of chlorinated and brominated Al compds. to give a composition having 1:90-50:50 Br-Cl mol ratio and 1:1.6-1:2 Mg-halogen mol ratio. Thus, a solution of 0.03 mol MgAl_2Et_8 [15415-18-6] in 150 mL heptane was treated with a solution containing 0.056 mol Et_2AlCl [96-10-6] and 0.002 mol AlBr_3 in 50 mL heptane for 6 h under reflux to give a precipitate containing Mg 20 , Al 0.1, Cl 47.3, and Br 13.1%. The precipitate was treated with EtOBz in heptane at 70° for 4 h, then with TiCl_4 at 140° for 3 h to give a catalyst containing Mg 20.7, Al 0.05, Ti 0.8, Br 9.7 and Cl 53.2%. Polymn of propylene using the catalyst and Et_3Al [97-93-8] gave isotactic polypropylene [25085-53-4] with intrinsic viscosity 3.7 dL/g and isotacticity 97%. Polymer yield was 112,500 g polymer/g Ti.

IT 82404-69-1
RL: CAT (Catalyst use); USES (Uses)
(catalysts, for polymerization of propylene)

RN 82404-69-1 CAPLUS

CN Aluminate(2-), pentaethyl-, magnesium (1:1) (CA INDEX NAME)



● Mg²⁺

L21 ANSWER 6 OF 10 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 1975:479316 CAPLUS
DOCUMENT NUMBER: 83:79316
ORIGINAL REFERENCE NO.: 83:12459a,12462a
TITLE: Reaction of alkaline earth metals with organomercury compounds in the presence of aluminumtrialkyls and aluminumtriaryls
AUTHOR(S): Ivanov, L. L.; Zavizion, S. Ya.; Zakharkin, L. I.

10/923,271

CORPORATE SOURCE: Inst. Elementoorg. Soedin., Moscow, USSR

SOURCE: Zhurnal Obshchei Khimii (1975), 45(5),
1060-5

CODEN: ZOKHA4; ISSN: 0044-460X

DOCUMENT TYPE: Journal

LANGUAGE: Russian

AB The aluminum compds. $M(AlR_3R_1)_2$ ($R = Et, Pr, Ph$; $R_1 = Et, Pr, Ph, MeC_6H_4$; $M = Ca, Sr, Ba$) were prepared by the reaction of AlR_3 with HgR_1_2 and M , with or without solvents (Et_2O , THF, Me_3N etc.). In the presence of solvents, the solvated products $M(AlR_3R_1)_2 \cdot nL$ ($n = 2, 3, 4, 6$; $L =$ solvent) were formed.

IT 56413-54-8P

RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of)

RN 56413-54-8 CAPLUS

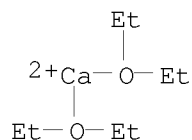
CN Calcium(2+), bis[1,1'-oxybis[ethane]]-, bis[(T-4)-tetraethylaluminate(1-)]
(9CI) (CA INDEX NAME)

CM 1

CRN 56413-53-7

CMF C8 H20 Ca O2

CCI CCS

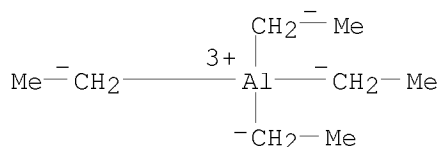


CM 2

CRN 14913-44-1

CMF C8 H20 Al

CCI CCS



L21 ANSWER 7 OF 10 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 1970:132840 CAPLUS

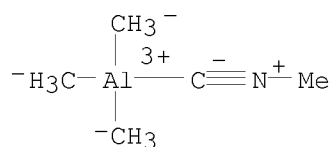
DOCUMENT NUMBER: 72:132840

ORIGINAL REFERENCE NO.: 72:23791a

TITLE: Reactions of methyl isocyanide with aluminum
compounds

AUTHOR(S): Meller, Anton; Batka, H.

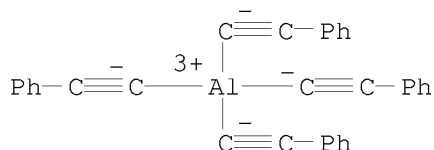
CORPORATE SOURCE: Inst. Anorg. Chem., Tech. Hochsch. Wien, Vienna, Austria
 SOURCE: Monatsh. Chem. (1970), 101(2), 627-8
 CODEN: MOCHAP
 DOCUMENT TYPE: Journal
 LANGUAGE: German
 GI For diagram(s), see printed CA Issue.
 AB Me₃Al.C.tplbond.NMe was obtained by treatment of Me₃Al with MeN.tplbond.C. It ignited spontaneously in both air and water. Treatment of AlCl₃ with MeN.tplbond.C gave 20% (C₆H₉AlCl₃N₃)₂ which had a cyclic structure (I).
 IT 27681-26-1P
 RL: SPN (Synthetic preparation); PREP (Preparation)
 (preparation of)
 RN 27681-26-1 CAPLUS
 CN Aluminum, [(isocyano-κC)methane]trimethyl-, (T-4)- (9CI) (CA INDEX NAME)



L21 ANSWER 8 OF 10 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 1965:498485 CAPLUS
 DOCUMENT NUMBER: 63:98485
 ORIGINAL REFERENCE NO.: 63:18132g-h,18133a
 TITLE: Reactions of organoaluminum compounds with acyl peroxides and anhydrides
 AUTHOR(S): Razuvaev, G. A.; Stepovik, L. P.
 CORPORATE SOURCE: State Univ., Gorki
 SOURCE: Zhurnal Obshchei Khimii (1965), 35(9), 1672-6
 CODEN: ZOKHA4; ISSN: 0044-460X
 DOCUMENT TYPE: Journal
 LANGUAGE: Russian
 AB A mixture of (iso-PrO)₃Al and 1 mole m-O₂NC₆H₄CO₂Ac in C₆H₆ under N gave, after evaporation and treatment with aqueous KOH, .apprx.10-12% AcOH, but m-O₂NC₆H₄CO₂CMe:CH₂ was not isolated. Similarly ClCH₂CO₂Ac gave ClCH₂CO₂H; o-O₂NC₆H₄CO₂Ac gave AcOH (4%) in 10-15 min. EtAl(OEt)₂ and 1 mole Bz₂O₂ in C₆H₆ under N gave in 3-4 days AcH, BzOEt, and BzOAl(OEt)₂; similar treatment with BzO₂Ac gave AcOH. The reaction of (iso-PrO)₃Al with mixed acid anhydrides gave the alkoxy-Al salts of the stronger acid and an ester of the weaker acid. (EtO)₂AlEt and acyl peroxides gave esters and alkoxy-Al salts of carboxylic acids. Both reactions appear to proceed through a complex formed at the Al atom with the O bridge of the anhydrides or 2 O atoms of the peroxides.
 IT 68446-25-3P, Sodium tetrakis(phenylethynyl)aluminate
 700798-30-7P, Aluminate, tetrakis(phenylethynyl)-
 744953-02-4P, Aluminate, tetra-1-hexynyl-
 RL: PREP (Preparation)
 (preparation of)
 RN 68446-25-3 CAPLUS

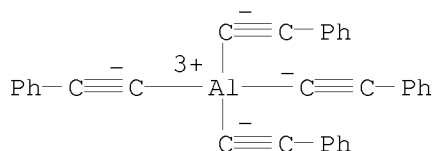
10/923,271

CN Aluminate(1-), tetrakis(phenylethynyl)-, sodium, (T-4)- (9CI) (CA INDEX NAME)



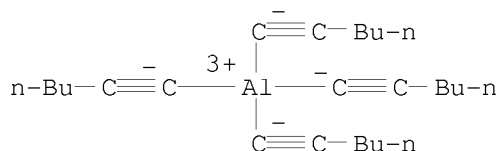
RN 700798-30-7 CAPLUS

CN Aluminate(1-), tetrakis(phenylethynyl)-, (T-4)- (9CI) (CA INDEX NAME)



RN 744953-02-4 CAPLUS

CN Aluminate(1-), tetra-1-hexynyl-, (T-4)- (9CI) (CA INDEX NAME)



L21 ANSWER 9 OF 10 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 1963:448453 CAPLUS

DOCUMENT NUMBER: 59:48453

ORIGINAL REFERENCE NO.: 59:8772e-g

TITLE: Synthesis of complex aluminum acetylides
MAl(C.tplbond.CR)₄, where M = Li, Na, or K, and their
reactions with carbonyl compds.

AUTHOR(S): Zakharkin, L. I.; Gavrilenko, V. V.

SOURCE: Izvestiya Akademii Nauk SSSR, Seriya Khimicheskaya (1963), (6), 1146-7

CODEN: IASKA6; ISSN: 0002-3353

DOCUMENT TYPE: Journal

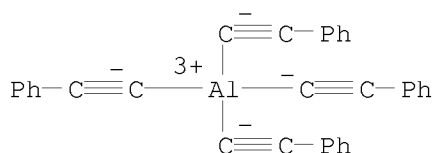
LANGUAGE: Unavailable

AB The reaction MAlH₄ + 4HC.tplbond.CR → MAl(C.tplbond.CR)₄ + 4H₂

(where R = alkyl, aryl, or H) can be carried out with NaAlH₄ or KAlH₄ as

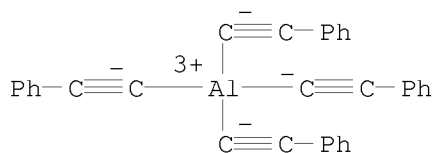
well as with LiAlH_4 . With $M = \text{Li}$ or Na , the reaction took place readily in tetrahydrofuran, while diglyme was the best solvent for the reaction with Li compds. Carbonyl compds. also reacted with the complex Al acetylides: $\text{MAl}(\text{C.tplbond.CR})_4 + :\text{CO} \rightarrow :\text{C}(\text{OH})\text{C.tplbond.CR}$. Thus, $\text{MAl}(\text{C.tplbond.CPh})_4$ ($M = \text{Li}, \text{Na}, \text{K}$) and PhCHO formed $\text{PhC.tplbond.CCH}(\text{OH})\text{Ph}$ (70-80% yield); $\text{NaAl}(\text{C.tplbond.CBu})_4$ and PrCHO formed $\text{BuC.tplbond.CCH}(\text{OH})\text{Pr}$ (70%); $\text{NaAl}(\text{C.tplbond.CBu})_4$ and crotonaldehyde formed $\text{MeCH:CHCH}(\text{OH})\text{C.tplbond.CBu}$ (80%); while $\text{NaAl}(\text{C.tplbond.CH})_4$ and butyraldehyde, PhCHO , or phenylacetone formed the corresponding acetylenic alcs. with yields of 40-50%,. At elevated temps., carboxylic acids could be prepared with good yields by the reaction $\text{MAl}(\text{C.tplbond.CR})_4 + \text{CO}_2 \rightarrow \text{RC.tplbond.CCO}_2\text{H}$. On passing CO_2 through a solution of $\text{NaAl}(\text{C.tplbond.CPh})_4$ in diglyme at $120-50^\circ$, 60% phenylpropionic acid was obtained.

IT 4015-69-4P, Lithium tetrakis(phenylethynyl)aluminate
 68446-25-3P, Sodium tetrakis(phenylethynyl)aluminate
 700798-30-7P, Aluminate, tetrakis(phenylethynyl)-
 744953-02-4P, Aluminate, tetra-1-hexynyl-
 RL: PREP (Preparation)
 (preparation of)
 RN 4015-69-4 CAPLUS
 CN Aluminate(1-), tetrakis(phenylethynyl)-, lithium, (T-4)- (9CI) (CA INDEX NAME)



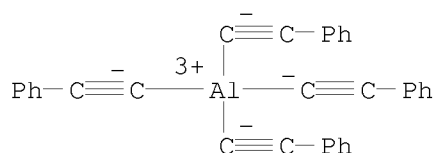
● Li^+

RN 68446-25-3 CAPLUS
 CN Aluminate(1-), tetrakis(phenylethynyl)-, sodium, (T-4)- (9CI) (CA INDEX NAME)



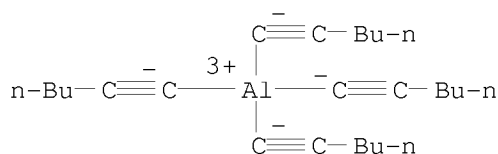
● Na^+

RN 700798-30-7 CAPLUS
 CN Aluminate(1-), tetrakis(phenylethynyl)-, (T-4)- (9CI) (CA INDEX NAME)



RN 744953-02-4 CAPLUS

CN Aluminate(1-), tetra-1-hexynyl-, (T-4)- (9CI) (CA INDEX NAME)



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ACCESSION NUMBER: 1963:53473 CAPLUS

DOCUMENT NUMBER: 58:53473

ORIGINAL REFERENCE NO.: 58:9135h,9136a-b

TITLE: Organometallic reactions

INVENTOR(S): Kobetz, Paul; Pinkerton, Richard C.

PATENT ASSIGNEE(S): Ethyl Corp.

SOURCE: 4 pp.

DOCUMENT TYPE: Patent

LANGUAGE: Unavailable

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 3068261		19621211	US 1960-5593	19600201 <--
PRIORITY APPLN. INFO.:			US	19600201

AB Alkali metal Al hydrocarbon complexes (I) are prepared by treating the corresponding B complexes with an Al trihydrocarbon compound. Thus, a stirred mixture of NaBEt₄ (II) 150 (1 mole) and AlEt₃ (III) 228 parts (2 moles) is heated to 125° to give volatile BEt₃ (IV) and a residue consisting of equimolar proportions of III and NaAlEt₄ (V). By cooling the mixture to room temperature, V is crystallized as a readily filterable solid.

Similarly are prepared (B reactant, moles, Al reactant, moles, I product, and B trihydrocarbon product given): LiBEt₄, 1, III, 3, LiAlEt₄, IV; II, 1, AlMe₃, 1, NaAlMe₃Et, IV; II, 1, AlMe₃, 3, V, BMe₃; NaBPr₄, 1, III, 2, NaAlEt₃Pr, BEt₃-iso Pr compds.; NaBPh₄, 1, Al(iso-Bu)₃, 1, mixture of NaAl-iso-Bu-Ph compds., BPh₃ + B(iso-Bu)₃; KBet₄, 1, AlPr₃, 1, KAlEtPr₃, IV; NaB(CH₂Ph)₄, 1, AlPh₃, 1, NaAlPh₃(CH₂Ph), B(CH₂Ph)₃. The reaction can be utilized for the selective separation of organometallic mixts. which include Al trihydrocarbon compds. as a component. Thus, a single phase liquid mixture of 42% PbEt₄ (VI) and 58% III 1000 (the mixture also containing

a

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fraction of 1% of a thermal stabilizer for VI) is treated with II 760 parts at 100° with vigorous agitation to give vaporized IV and a mixture of immiscible V and VI. I are useful as alkylating agents in producing organo-metallic compds. of other metals and as electrolyte components for electrolytic processes. The B trihydrocarbon materials released in the process are valuable as components of high energy fuel compns.

IT 701193-48-8P, Aluminate, ethyltrimethyl-

RL: PREP (Preparation)

(preparation of)

RN 701193-48-8 CAPLUS

CN Aluminate(1-), ethyltrimethyl-, (T-4)- (CA INDEX NAME)

